

DOCKET NO: 17302(HL)

PATENT

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicant: Forman et al)

Group Art Unit: 1617

Serial No.: 09/590,447)

Conf. No.: 1446)

Filed: June 9, 2000)

For: METHODS FOR)

MODULATING)

FXR RECEPTOR ACTIVITY)

Examiner: Hui, S.)

I hereby certify that this correspondence is being deposited with the United States Postal Service as First Class Mail in an envelope addressed to: Mail Stop Amendment-Fee, Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450 on:

Date of Deposit: 6/30/2003

Person making Deposit: Bonnie Ferguson

Signature: Bonnie Ferguson

Date of Signature: 6/30/2003

AMENDMENT

Commissioner for Patents
Alexandria, VA 22313-1450

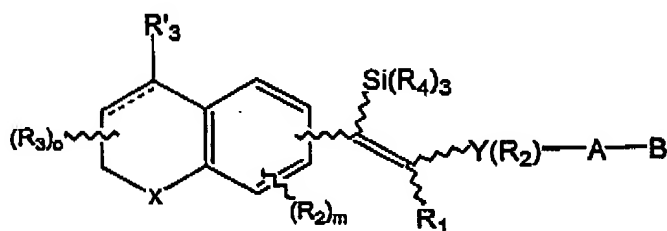
Dear Sir:

This communication is responsive to the Office Action received January 29, 2003. The Office Action has been carefully considered, and Applicants have the following comments.

Serial No. : 09/590,447
 Filed : June 9, 2000

AMENDMENTS AND STATUS OF CLAIMS

1. (Currently amended) A method of treating an FXR-mediated pathological condition in a mammal comprising the step of administering to a mammal in need thereof a pharmaceutically acceptable composition comprising a synthetic FXR ligand able to stimulate, block, or inhibit the activity of a mammalian FXR receptor, said synthetic FXR ligand comprising a compound of the formula



formula (3)

wherein the dashed line represents a bond or absence of a bond;

X is S, O, NR' where R' is H or alkyl of 1 to 6 carbons, or

X is $(C(R_1)_2)_n$ where R_1 is H or alkyl of 1 to 6 carbons, and n is an integer having the value of 0 or 1;

R_2 is hydrogen, lower alkyl of 1 to 6 carbons, F, Cl, Br, I, CF_3 , fluoro substituted alkyl of 1 to 6 carbons, OH, SH, alkoxy of 1 to 12 carbons, or alkylthio of 1 to 12 carbons, benzyloxy or $C_1 - C_{12}$ alkylbenzyloxy;

R_3 is hydrogen, lower alkyl of 1 to 6 carbons or F;

m is an integer having the value of 0 - 3;

o is an integer having the value of 0 - 4 when the dashed line represents absence of a bond, and 0 - 3 when the dashed line represents a bond;

$[R_3]$ R'_3 is hydrogen, lower alkyl of 1 to 6 carbons, F or $(R_{15})_r$ -phenyl, $(R_{15})_r$ -naphthyl, or $(R_{15})_r$ -heteroaryl where the heteroaryl group has 1 to 3 heteroatoms selected from the group consisting of O, S and N, r is an integer having the values of 0 - 5;

Serial No. : 09/590,447
Filed : June 9, 2000

R_4 is alkyl of 1 to 8 carbons, or phenyl;

Y is a phenyl or naphthyl group, or heteroaryl selected from a group consisting of pyridyl, thienyl, furyl, pyridazinyl, pyrimidinyl, pyrazinyl, thiazolyl, oxazolyl, imidazolyl and pyrazolyl, said phenyl and heteroaryl groups being optionally substituted with one or two R_2 groups;

R_{15} is independently H, F, Cl, Br, I, NO_2 , $N(R_8)_2$, $NH(R_8)$, COR_8 , $NR_8CON(R_8)_2$, OH , $OCOR_8$, OR_8 , CN, an alkyl group having 1 to 10 carbons, fluoro substituted alkyl group having 1 to 10 carbons, an alkenyl group having 1 to 10 carbons and 1 to 3 double bonds, alkynyl group having 1 to 10 carbons and 1 to 3 triple bonds, or a trialkylsilyl or trialkylsilyloxy group where the alkyl groups independently have 1 to 6 carbons;

A is $(CH_2)_q$ where q is 0-5, lower branched chain alkyl having 3-6 carbons, cycloalkyl having 3-6 carbons, alkenyl having 2-6 carbons and 1 or 2 double bonds, alkynyl having 2-6 carbons and 1 or 2 triple bonds;

B is hydrogen, $COOH$, NO_2 , $P(O)(OH)_2$, $P(O)(OH)OR_8$, $P(O)(OR_8)_2$, SO_2OH , $SO_2(OR_8)$, $COOR_8$, $CONR_9R_{10}$, $-CH_2OH$, CH_2OR_{11} , CH_2OCOR_{11} , CHO , $CH(OR_{12})_2$, $CHOR_{13}O$, $-COR_7$, $CR_7(OR_{12})_2$, $CR_7OR_{13}O$, or tri-lower alkylsilyl, where R_7 is an alkyl, cycloalkyl or alkenyl group containing 1 to 5 carbons, R_8 is an alkyl group of 1 to 10 carbons or trimethylsilylalkyl where the alkyl group has 1 to 10 carbons, or a cycloalkyl group of 5 to 10 carbons, or R_8 is phenyl or lower alkylphenyl, R_9 and R_{10} independently are hydrogen, an alkyl group of 1 to 10 carbons, or a cycloalkyl group of 5-10 carbons, or phenyl or lower alkylphenyl, R_{11} is lower alkyl, phenyl or lower alkylphenyl, R_{12} is lower alkyl, and R_{13} is divalent alkyl radical of 2-5 carbons, or a pharmaceutically acceptable salt of said compound.

2. (Original) A method in accordance with Claim 1 where X is $(C(R_1)_2)_n$ and n is 1.
3. (Original) A method in accordance with Claim 1 where X is S.
4. (Original) A method in accordance with Claim 1 where X is O.
5. (Original) A method in accordance with Claim 1 where X is NR.
6. (Original) A method in accordance with Claim 1 where Y is phenyl.
7. (Original) A method in accordance with Claim 1 where Y is thienyl.

Serial No. : 09/590,447
Filed : June 9, 2000

8. (Original) A method in accordance with Claim 1 wherein said compound has a structure selected from formulas (1) and (2).

9. (Original) A method in accordance with Claim 8 wherein said compound has a structure of formula (1) where the dashed line represents absence of a bond.

10. (Original) A method in accordance with Claim 8 wherein said compound has a structure of formula (1) where the dashed line represents a bond.

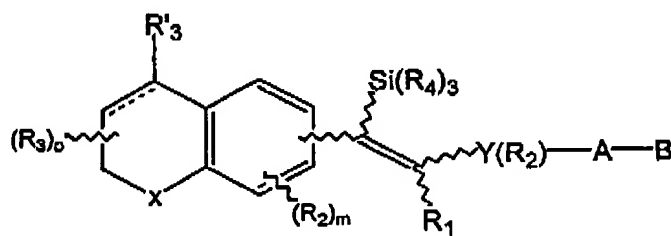
11. (Original) A method in accordance with Claim 1 wherein said compound has a structure selected from formulas (3) and (4).

12. (Original) A method in accordance with Claim 11 wherein said compound has a structure of formula (3) where the dashed line represents absence of a bond.

13. (Original) A method in accordance with Claim 11 wherein said compound has a structure of formula (3) where the dashed line represents a bond.

14-30 (Withdrawn)

31. (Previously amended) A method of treating a hypercholesterolemic mammal comprising the steps: administering to a mammal in need thereof a pharmaceutically acceptable composition comprising an FXR antagonist having the following formula



formula (3)

wherein the dashed line represents a bond or absence of a bond;

X is S, O, NR' where R' is H or alkyl of 1 to 6 carbons, or

X is $(C(R_1)_2)_n$ where R_1 is H or alkyl of 1 to 6 carbons, and n is an integer having the value of 0 or 1;

R₂ is hydrogen, lower alkyl of 1 to 6 carbons, F, Cl, Br, I, CF₃, fluoro substituted

Serial No. : 09/590,447
Filed : June 9, 2000

alkyl of 1 to 6 carbons, OH, SH, alkoxy of 1 to 12 carbons, or alkylthio of 1 to 12 carbons, benzyloxy or $C_1 - C_{12}$ alkylbenzyloxy;

R_3 is hydrogen, lower alkyl of 1 to 6 carbons or F;

m is an integer having the value of 0 - 3;

o is an integer having the value of 0 - 4 when the dashed line represents absence of a bond, and 0 - 3 when the dashed line represents a bond;

R'_3 is hydrogen, lower alkyl of 1 to 6 carbons, F or $(R_{15})_r$ -phenyl, $(R_{15})_r$ -naphthyl, or $(R_{15})_r$ -heteroaryl where the heteroaryl group has 1 to 3 heteroatoms selected from the group consisting of O, S and N, r is an integer having the values of 0 - 5;

R_4 is alkyl of 1 to 8 carbons, or phenyl;

Y is a phenyl or naphthyl group, or heteroaryl selected from a group consisting of pyridyl, thienyl, furyl, pyridazinyl, pyrimidinyl, pyrazinyl, thiazolyl, oxazolyl, imidazolyl and pyrazolyl, said phenyl and heteroaryl groups being optionally substituted with one or two R_2 groups;

R_{15} is independently H, F, Cl, Br, I, NO_2 , $N(R_8)_2$, $NH(R_8)$, COR_8 , $NR_8CON(R_8)_2$, OH, $OCOR_8$, OR_8 , CN, an alkyl group having 1 to 10 carbons, fluoro substituted alkyl group having 1 to 10 carbons, an alkenyl group having 1 to 10 carbons and 1 to 3 double bonds, alkynyl group having 1 to 10 carbons and 1 to 3 triple bonds, or a trialkylsilyl or trialkylsilyloxy group where the alkyl groups independently have 1 to 6 carbons;

A is $(CH_2)_q$ where q is 0-5, lower branched chain alkyl having 3-6 carbons, cycloalkyl having 3-6 carbons, alkenyl having 2-6 carbons and 1 or 2 double bonds, alkynyl having 2-6 carbons and 1 or 2 triple bonds;

B is hydrogen, COOH, NO_2 , $P(O)(OH)_2$, $P(O)(OH)OR_8$, $P(O)(OR_8)_2$, SO_2OH , $SO_2(OR_8)$, $COOR_8$, $CONR_9R_{10}$, $-CH_2OH$, CH_2OR_{11} , CH_2OCOR_{11} , CHO, $CH(OR_{12})_2$, $CHOR_{13}O$, $-COR_7$, $CR_7(OR_{12})_2$, $CR_7OR_{13}O$, or tri-lower alkylsilyl, where R_7 is an alkyl, cycloalkyl or alkenyl group containing 1 to 5 carbons, R_8 is an alkyl group of 1 to 10 carbons or trimethylsilylalkyl where the alkyl group has 1 to 10 carbons, or a cycloalkyl group of 5 to 10 carbons, or R_8 is phenyl or lower alkylphenyl, R_9 and R_{10} independently are hydrogen, an alkyl group of 1 to 10 carbons, or a cycloalkyl group of 5-10 carbons, or phenyl or lower alkylphenyl, R_{11} is lower alkyl, phenyl or lower alkylphenyl, R_{12} is lower alkyl, and R_{13} is divalent alkyl radical of 2-5 carbons, or a pharmaceutically acceptable salt of said compound.

Serial No. : 09/590,447
Filed : June 9, 2000

32. (Currently amended) A method of treating an FXR-mediated pathological condition in a mammal comprising the step of providing to said mammal a pharmaceutically acceptable composition comprising a synthetic FXR ligand able to stimulate, block, or inhibit the activity of a mammalian FXR receptor.

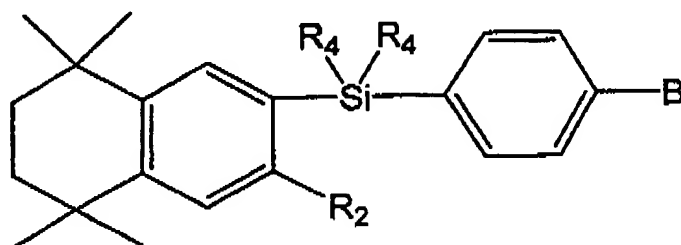
33. (Original) The method of claim 32 wherein said pathological condition comprises hypercholesterolemia.

34. (Original) The method of claim 32 wherein said pathological condition comprises hypocholesterolemia.

35. (Original) The method of claim 32 wherein said pathological condition is characterized by the overproduction of bile acids.

36. (Original) The method of claim 32 wherein said pathological condition is characterized by the underproduction of bile acids.

37. (Currently amended) A method of treating an FXR-mediated pathological condition in a mammal comprising the step of administering to a mammal in need thereof a pharmaceutically acceptable composition comprising a synthetic FXR ligand able to stimulate, block, or inhibit the activity of a mammalian FXR receptor, said synthetic FXR ligand having the formula



wherein R₂ is H or lower alkyl, R₄ is lower alkyl of 1 to 8 carbons and B is CH₂OH or COOR₈ where R₈ is H or ethyl.

38. (Original) A method in accordance with Claim 31 where R₂ is H and R₄ is ethyl.

39. (Original) A method in accordance with Claim 32 where B is CH₂OH.

40. (Original) A method in accordance with Claim 33 where B is COOR₈.

Serial No. : 09/590,447
Filed : June 9, 2000

REMARKS

Applicants thank the Examiner for the helpful suggestions contained in the January 29, 2003 Office Action, and for indicating that the claims, amended as above, are allowable. The claim Amendments made herein render moot the rejections pursuant to 35 USC 112(1) and 35 USC 112(2)

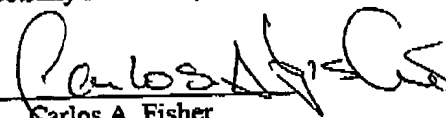
Therefore the claims appear to be in condition for allowance. Please use Deposit Account 01-0885 for the payment of the extension fees or any other fees due in connection with the current response.

Respectfully submitted,

Dated:

6/30/03

By:



Carlos A. Fisher
Registration No. 36,510
Attorney of Record

Allergan, Inc.
2525 Dupont Drive
Irvine, CA 92612
Telephone: 714-246-4920
Facsimile: 714-246-4249

COPY

MAIL STOP AMENDMENT-FEE

Rec'd in USPTO/PCT Office. Date Stamp and Return Card.

Date: 6/30/2003 Serial No.: 09/590,447; Conf.No. 1446

Title: METHODS FOR MODULATING FXR RECEPTOR.

Dkt. No.: 17302 (HL)

Enclosed Are:

- ☐ Specification # _____, Claims # _____, and Abstract # _____
- ☐ Drawings (_____ sheets)
_____ Formal _____ Informal
- ☐ Info. Disc. Statement
- ☐ Priority Documents # _____
- ☐ PTO 1449 W/References
- ☐ PCT Request (# pgs. _____)
- ☐ PCT Demand (# pgs. _____)
- ☐ PCT Response (# pgs. _____)
- ☐ PCT Amendment (# pgs. _____)

- ☐ Declaration, Power of Attorney
- ☐ Assignment & Cover Sheet
- ☒ Amendment (Final) (# pgs. 7)
- ☒ Certificate of Mailing
- ☐ Issue Fee Transmittal
- ☐ Transmittal Letter
- ☒ Extension of Time 2 months
- ☐ Express Mail No. _____

MAIL STOP AMENDMENT-FEE

Rec'd in USPTO/PCT Office. Date Stamp and Return Card.

Date: 6/30/2003 Serial No.: 09/590,447; Conf.No. 1446

Title: METHODS FOR MODULATING FXR RECEPTOR.

Dkt. No.: 17302 (HL)

Enclosed Are:

- ☐ Specification # _____, Claims # _____, and Abstract # _____
- ☐ Drawings (_____ sheets)
_____ Formal _____ Informal
- ☐ Info. Disc. Statement
- ☐ Priority Documents # _____
- ☐ PTO 1449 W/References
- ☐ PCT Request (# pgs. _____)
- ☐ PCT Demand (# pgs. _____)
- ☐ PCT Response (# pgs. _____)
- ☐ PCT Amendment (# pgs. _____)
- ☐ Declaration, Power of Attorney
- ☐ Assignment & Cover Sheet
- ☒ Amendment (Final) (# pgs. 7)
- ☒ Certificate of Mailing
- ☐ Issue Fee Transmittal
- ☐ Transmittal Letter
- ☒ Extension of Time 2 months
- ☒ Express Mail No. RECEIVED



LEGAL DEPARTMENT

2525 Dupont Drive
Irvine, California 92612

TO: Sam Hui

Fax No. 703-746-3123

FROM: CARLOS A. FISHER

Telephone-714-246-4920

FAX NO.: 714-246-4249

DATE: October 28, 2003

Pages being sent including this cover page:

cc:

Re: Privacy Gap Analysis

This fax has been sent to one number only- please copy ALL OTHER addressees in your location/country.

CONFIDENTIAL / ATTORNEY CLIENT PRIVILEGED COMMUNICATION

The information contained in this transmission is privileged and confidential. It is intended only for the use of the individual or entity named below. If the reader of this message is not the intended recipient or the employee or agent responsible for delivering the message to the intended recipient, you are hereby notified that any dissemination, distribution, or copying of this communication is strictly prohibited. If you have received this communication in error, please notify Allergan immediately by telephone and return the original message to us at the above-indicated address via regular U.S. mail. Thank you.

☒ ORIGINAL WILL NOT FOLLOW

☐ ORIGINAL WILL FOLLOW VIA:

- ☐ Regular Mail
- ☐ Overnight Delivery
- ☐ Hand Delivery
- ☐ Other

Attached is a copy of the Response as filed on July 31, 2003.



UNITED STATES PATENT AND TRADEMARK OFFICE

UNDER SECRETARY OF COMMERCE FOR INTELLECTUAL PROPERTY AND
DIRECTOR OF THE UNITED STATES PATENT AND TRADEMARK OFFICE
WASHINGTON, DC 20231
www.uspto.gov

RECEIVED

JUL 29 2003

LEGAL/PATENTS Paper No.

Notice of Non-Compliant Amendment (Voluntary Revised Practice)

The amendment filed July 2, 2003 under the voluntary revised amendment practice guidelines¹, published in the Official Gazette on February 25, 2003 (*Amendments in a Revised Format Now Permitted*, 1267 Off. Gazette 106), does not fully comply with minimal requirements of the voluntary practice. In order for the amendment to be entered, it must either (1) comply with the guidelines of the voluntary revised amendment practice (which practice invokes waivers of certain 37 CFR 1.121(a)-(d) requirements) or (2) comply with current 37 CFR 1.121 requirements.

THE FOLLOWING ITEM(S) IN APPLICANT'S AMENDMENT CAUSES THE AMENDMENT TO BE NON-COMPLIANT WITH THE VOLUNTARY REVISED AMENDMENT PRACTICE.

- ☐ 1. A complete listing of all of the claims is not present in the amendment paper.
- ☐ 2. The listing of claims does not include the text of all claims currently under examination.
- ☐ 3. The claims of this amendment paper have not been presented in ascending numerical order.
- ☒ 4. Each claim has not been provided with a status identifier, and, as such, the individual status of each claim cannot be determined.
- ☒ 5. Other: Claims 14-30 have the wrong status identifier

LIE: Check one of the following boxes:

- ☐ **PRELIMINARY AMENDMENT:** Applicant is given ONE MONTH from the mail date of this letter to re-submit the amendment in compliance with either the guidelines of the revised amendment practice or current 37 CFR 1.121. Failure to comply with either the current 37 CFR 1.121 practice or with the voluntary practice will result in non-entry of the amendment and examination on the merits will commence without entry of the originally proposed preliminary amendment. This notice is not an action under 35 U.S.C. 132, and this ONE MONTH time limit is not extendable.

- ☒ **AMENDMENT AFTER NON-FINAL ACTION:** Since the above-mentioned reply appears to be a *bona fide* response, applicant is given a TIME PERIOD of ONE MONTH from the mailing of this notice within which to re-submit an amendment which complies with either the voluntary practice guidelines or current 37 CFR 1.121 in order to avoid abandonment. EXTENSIONS OF THIS TIME PERIOD ARE AVAILABLE UNDER 37 CFR 1.136(a).

Brenda Gray
Team Leader

¹ For further explanation of the guidelines of the revised amendment format, please see the posted notice and sample amendment format at:
<http://www.uspto.gov/web/offices/pac/dapp/npla/preognotice/officeflyer.pdf> and
<http://www.uspto.gov/web/offices/pac/dapp/npla/preognotice/formatrevandtrac.pdf>

MAIL STOP-REPLY-NON-COMPLIANCE

Rec'd in USPTO/PCT Office. Date Stamp and Return Card.

Date: 7/31/2003

Serial No.: 09/590,447; Conf. No. 1446

Title: METHODS FOR MODULATING FXR RECEPTOR

Dkt. No.: 17302(HL)

Enclosed Are:

- | | |
|--|---|
| <input type="checkbox"/> Specification # _____, Claims # _____, and Abstract # _____ | <input type="checkbox"/> Declaration, Power of Attorney |
| <input type="checkbox"/> Drawings (____ sheets)
____ Formal ____ Informal | <input type="checkbox"/> Assignment & Cover Sheet |
| <input type="checkbox"/> Info. Disc. Statement | <input checked="" type="checkbox"/> Amendment (Elnat) (# pgs. ____) |
| <input type="checkbox"/> Priority Documents # _____ | <input checked="" type="checkbox"/> Certificate of Mailing |
| <input type="checkbox"/> PTO 1449 W/References | <input type="checkbox"/> Issue Fee Transmittal |
| <input type="checkbox"/> PCT Request (# pgs. ____) | <input type="checkbox"/> Transmittal Letter |
| <input type="checkbox"/> PCT Demand (# pgs. ____) | <input type="checkbox"/> Extension of Time |
| <input type="checkbox"/> PCT Response (# pgs. ____) | <input type="checkbox"/> Express Mail No. _____ |
| <input type="checkbox"/> PCT Amendment (# pgs. ____) | <input checked="" type="checkbox"/> COPY OF 6/29/03 Amend. |
| | <input checked="" type="checkbox"/> Corrected Claims |
| | <input checked="" type="checkbox"/> NON-COMPLIANCE NOTICE |

MAIL STOP-REPLY-NON-COMPLIANCE

Rec'd in USPTO/PCT Office. Date Stamp and Return Card.

Date: 7/31/2003

Serial No.: 09/590,447; Conf. No. 1446

Title: METHODS FOR MODULATING FXR RECEPTOR

Dkt. No.: 17302(HL)

Enclosed Are:

- | | |
|--|--|
| <input type="checkbox"/> Specification # _____, Claims # _____, and Abstract # _____ | <input type="checkbox"/> Declaration, Power of Attorney |
| <input type="checkbox"/> Drawings (____ sheets)
____ Formal ____ Informal | <input type="checkbox"/> Assignment & Cover Sheet |
| <input type="checkbox"/> Info. Disc. Statement | <input checked="" type="checkbox"/> Amendment (Elnat) |
| <input type="checkbox"/> Priority Documents # _____ | <input type="checkbox"/> Certificate of Mailing |
| <input type="checkbox"/> PTO 1449 W/References | <input type="checkbox"/> Issue Fee Transmittal |
| <input type="checkbox"/> PCT Request (# pgs. ____) | <input type="checkbox"/> Transmittal Letter |
| <input type="checkbox"/> PCT Demand (# pgs. ____) | <input type="checkbox"/> Extension of Time |
| <input type="checkbox"/> PCT Response (# pgs. ____) | <input type="checkbox"/> Express Mail No. _____ |
| <input type="checkbox"/> PCT Amendment (# pgs. ____) | <input checked="" type="checkbox"/> COPY OF 6/29/03 Amend. |
| | <input checked="" type="checkbox"/> Corrected Claims |
| | <input checked="" type="checkbox"/> NON-COMPLIANCE NOTICE |

